

## EXHIBIT D

AFB fatty

or vegetable oil (corn, sunflower, peanut, palm, canola)

liquefied animal fat, preparations  
Lard, butter, fish oil, lard, tallow  
for pet foods, ammonium sulfide?

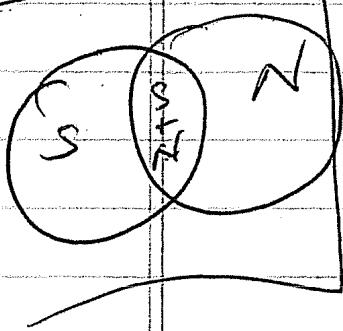
sulfur - donating reagent

eg, ① salt, such as sulfide (Na, K, Ca, etc)  
② cysteine, methionine  
③ cystine (2 S's linked together)

other S-containing A's

③ short peptides, such as glutathione  
④ elemental sulfur ( $S_8$ , yellow powder)  
recombine w/ fat or fat residues  
⑤ "sulfide liquor" from industrial (eg, paper, molasses, corn steep liquor)  
initial goal - add S to a fatty carrier that

① won't let powder settle out -  
powders typically don't mix solubly  
in fat  
easier to handle; sticks great



~~or~~ ~~or~~

Ammonium sulfide - could work, but  
odor problems likely - smel. on final  
food, if any Am remains

## N-donating reagents

① ~~Organic~~ micro-org such as  
yeast - est. 10%  $^{13}\text{N}$  by dry weight  
fungal mycelia (strands)

food-grade bacteria, known high N content

② basic A A's

arginine ( $\text{H}_2\text{N}-\text{CH}_2-\text{CH}_2-\text{NH}_2$ )

③ urea -

④ nucleotides -

more eff. than yeast, but possible

⑤ guanidino groups

⑥ heterocyclics -

but only if reactive, ready to donate  
up lot of cooking costs

~~Pektin~~

~~Pektin~~

distiller/brewers/bakers yeast

torula yeast - used to break down

"sulfide lignin" from paper-making

Two main ways

- ① low heat =  $< 98^{\circ}\text{C}$   
(anything under boiling).
- ② under pressure

reflux - tends to be lower losses

typical, if pressure used

~~120 to 200~~  $^{\circ}\text{C}$

10 psig to ~~100~~ psig (gen. by heat  
15 min to 1 hr; with ramps  
temperature

typical, if reflux used

~~90 to 98~~  $^{\circ}\text{C}$

ambient pressure, but closed vessel

1 to 6 hours

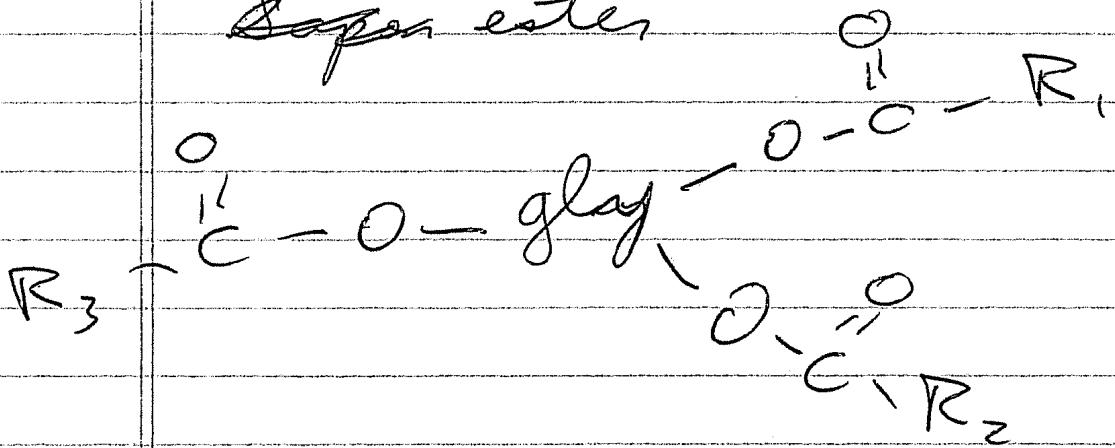
condensate ~~of~~ returned to <sup>cooking</sup> main vessel

Parr vessel = pressure cooker  
pressure, programmable

triglycerides [veg. oils - triglycerides]  
but diff FA's  
fat = combination of fatty acids  
+ glycerol (3 carbons), alcohol, triol

glycogen = mostly muscle, liver, in fat  
for quick breakdown

in fat -  
covalent bonds betw. FA's + glycerol  
sapon ester



where  $\text{R}_1 - \text{R}_3$  are FA's

saponification - breaks the ester bond  
can be complete or partial  
will regenerate  $\text{COOH}$  ( $\text{COO}^-$ ) group  
on FA's

the N + S will react w/ C's where bonds were broken in HCC chains - the heat breaks those bonds

goal - get smaller pieces/chunks from HCs + to c'lubinate with N and/or S attached to small pieces

lams -  
liquefied triglyceride prep -  
can be saponif before N/S addition

one good case of S added, w/o N source

S tends to give roasted flavor, "savory"  
dogs tend to prefer cats less so, but they like fat-derived  
preps